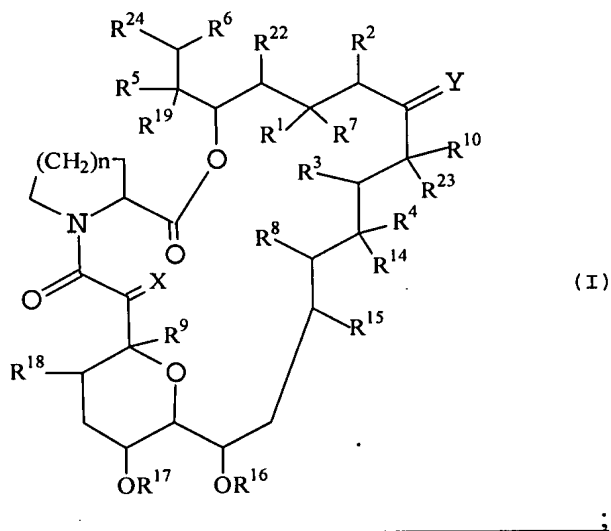


IN THE CLAIMS

Please amend the claims as follows:

Claim 1 (Currently Amended): A method of treating a human patient suffering from ~~an ocular allergy~~ allergic conjunctivitis, comprising:

administering to said patient an ophthalmic composition comprising from about ~~0.01% to about 0.1%~~ 0.03% to about 0.06% of a macrolide compound having the following formula (I), or a pharmaceutically acceptable salt thereof:



wherein adjacent pairs of R<sup>1</sup> and R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup>, and R<sup>5</sup> and R<sup>6</sup> each independently:

a) consist of two adjacent hydrogen atoms, wherein R<sup>2</sup> is optionally alkyl, or

b) form another bond optionally between carbon atoms binding with the members of

said pairs;

R<sup>7</sup> is a hydrogen atom, hydroxy, alkyloxy or protected hydroxy, or may form oxo with R<sup>1</sup>;

R<sup>8</sup> and R<sup>9</sup> each independently show hydrogen atom or hydroxy;

R<sup>10</sup> is a hydrogen atom, alkyl, alkyl substituted by one or more hydroxy, alkenyl, alkenyl substituted by one or more hydroxy or alkyl substituted by oxo;

X is oxo, (hydrogen atom, hydroxy), (hydrogen atom, hydrogen atom), or a group of the formula -CH<sub>2</sub>O-;

Y is oxo, (hydrogen atom, hydroxy), (hydrogen atom, hydrogen atom), or a group of the formula N-NR<sup>11</sup>R<sup>12</sup> or N-OR<sup>13</sup>;

R<sup>11</sup> and R<sup>12</sup> each independently are a hydrogen atom, alkyl, aryl or tosyl;

R<sup>13</sup>, R<sup>14</sup>, R<sup>15</sup>, R<sup>16</sup>, R<sup>17</sup>, R<sup>18</sup>, R<sup>19</sup>, R<sup>22</sup> and R<sup>23</sup> each independently are a hydrogen atom or alkyl;

R<sup>24</sup> is an optionally substituted ring that may contain one or more hetero atom(s); and  
n is 1 or 2.

Claims 2-3 (Cancelled)

Claim 4 (Previously Presented): A method according to claim 1, wherein said ophthalmic composition comprises about 0.03% of said macrolide compound.

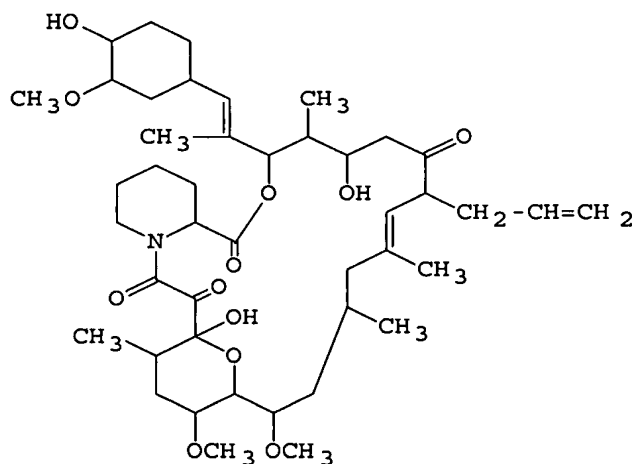
Claim 5 (Previously Presented): A method according to claim 1, wherein said macrolide compound is FK506.

Claim 6 (Previously Presented): A method according to claim 1, wherein said ophthalmic composition is an eye drop.

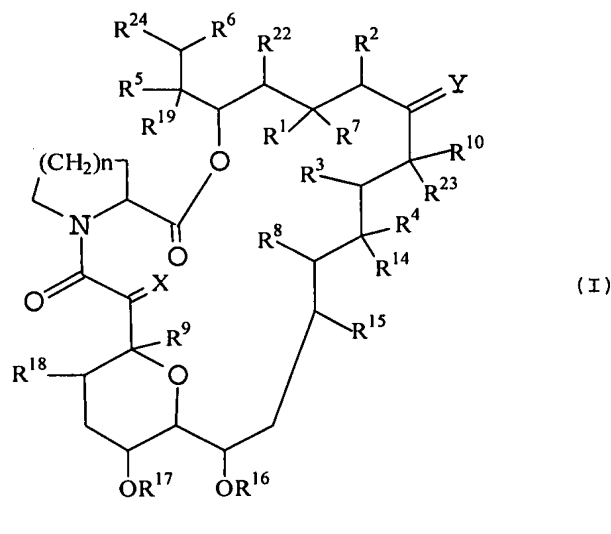
Claim 7 (Previously Presented): A method according to claim 6, wherein said eye drop further comprises polyvinyl alcohol.

Claim 9 (Original): A method according to claim 8, wherein said eye drop is administered from about one to about 4 times per day.

Claim 11 (Currently Amended): [[A]] The method according to claim [[10]] 1, wherein said macrolide compound has the following structure:



Claim 12 (Currently Amended): An ophthalmic composition for treatment of ocular allergy allergic conjunctivitis comprising from about 0.01% to about 0.1% 0.03 to about 0.06% of a macrolide compound having the following formula (I), or a pharmaceutically acceptable salt thereof:



wherein adjacent pairs of R<sup>1</sup> and R<sup>2</sup>, R<sup>3</sup> and R<sup>4</sup>, and R<sup>5</sup> and R<sup>6</sup> each independently  
a) consist of two adjacent hydrogen atoms, wherein R<sup>2</sup> is optionally alkyl, or  
b) form another bond optionally between carbon atoms binding with the members of  
said pairs;

R<sup>7</sup> is a hydrogen atom, hydroxy, alkyloxy or protected hydroxy, or may form oxo  
with R<sup>1</sup>;

R<sup>8</sup> and R<sup>9</sup> each independently are a hydrogen atom or hydroxy;

R<sup>10</sup> is hydrogen atom, alkyl, alkyl substituted by one or more hydroxy, alkenyl,  
alkenyl substituted by one or more hydroxy or alkyl substituted by oxo;

X is oxo, (hydrogen atom, hydroxy), (hydrogen atom, hydrogen atom), or a group of the formula -CH<sub>2</sub>O-;

Y is oxo, (hydrogen atom, hydroxy), (hydrogen atom, hydrogen atom), or a group of the formula N-NR<sup>11</sup>R<sup>12</sup> or N-OR<sup>13</sup>;

R<sup>11</sup> and R<sup>12</sup> each independently are a hydrogen atom, alkyl, aryl or tosyl;

R<sup>13</sup>, R<sup>14</sup>, R<sup>15</sup>, R<sup>16</sup>, R<sup>17</sup>, R<sup>18</sup>, R<sup>19</sup>, R<sup>22</sup> and R<sup>23</sup> each independently show hydrogen atom or alkyl;

R<sup>24</sup> is an optionally substituted ring that may contain one or more hetero atom(s); and  
n is 1 or 2.

Claims 13-14 (Cancelled)

Claim 15 (Previously Presented): An ophthalmic composition according to claim 12, comprising about 0.03% of said macrolide compound.

Claim 16 (Previously Presented): An ophthalmic composition according to claim 12, wherein said macrolide compound is FK506.

Claim 17 (Previously Presented): An ophthalmic composition according to claim 12, which is an eye drop.

Claim 18 (Previously Presented): An ophthalmic composition according to claim 17, wherein said eye drop further comprises polyvinyl alcohol.

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Claim 19 (Previously Presented): An ophthalmic composition according to claim 18, wherein said eye drop comprises about 0.03% of said macrolide compound.

Claim 20 (Currently Amended): An ophthalmic composition according to claim 19, wherein said eye drop is formulated in a form suitable for administration ~~administered~~ from about one to about 4 times per day.

Claims 21-33 (Canceled)

Claim 34 (Previously Presented): A commercial package comprising the ophthalmic composition of claim 12 and a written matter associated therewith, the written matter stating that the composition can or should be used for allergic conjunctivitis.